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Abstract

In view of the extensive use of hexagonal boron nitride (hBN) in 2D material electronics, it becomes important to refine its dielectric characterization in terms of low-field permittivity and high-field strength and conductivity up to the breakdown voltage. The present study aims at filling this gap using DC and RF transport in two Au-hBN-Au capacitor series of variable thickness in the 10–100 nm range, made of large high-pressure, high-temperature (HPHT) crystals and a polymer derivative ceramics (PDC) crystals. We deduce an out-of-plane low field dielectric constant $\epsilon_{\parallel} = 3.4 \pm 0.2$ consistent with the theoretical prediction of Ohba *et al*, that narrows down the generally accepted window $\epsilon_{\parallel} = 3-4$. The DC-current leakage at high-field is found to obey the Frenkel-Pool law for thermally-activated trap-assisted electron transport with a dynamic dielectric constant $\epsilon_{\parallel} \simeq 3.1$ and a trap energy $\Phi_B \simeq 1.3$ eV, that is comparable with standard technologically relevant dielectrics.

1. Introduction

Hexagonal boron nitride (hBN) is a van der Waals crystal insulator introduced in graphene electronics a decade ago [1] and since then extensively used as encapsulant [2], tunnel barrier [3–5], or gate dielectric in 2D material electronics [6, 7]. In view of its technological relevance, it is important to improve its characterization both in terms of low-field permittivity and high-field dielectric conductivity and breakdown field. Accepted values for the DC dielectric permittivity constant lie in the broad range $\epsilon^{\parallel} = 3-4$ [1, 8–10], whereas breakdown fields are more scattered, with $E_{BD} = 4-10$ MV cm⁻¹ [9, 11], depending on material quality and breakdown-field definition criterion. The purpose of the present study, which is based on DC and RF transport in Au-hBN-Au capacitors made of two types of hBN crystals, is to narrow down the uncertainty in permittivity, to shed light onto the dielectric breakdown mechanism, and to use these characterizations to benchmark the two hBN crystal sources.

We have used two series of Au-hBN-Au capacitors made of large exfoliated hBN crystals. Crystals are grown either under high-pressure high-temperature (HPHT samples) as described in [12], or with a polymer derivative ceramics (PDC) route described in [13]. Exfoliated hBN flakes, of thickness $d \sim 10-100$ nm, are sandwiched between Au electrodes of lateral dimensions $L \times W = 10 \times 10$ μ m. A significant fraction of samples (17 HPHTs and 11 PDCs of the total 41 capacitors) follows the parallel-plate capacitance law with a dielectric constant

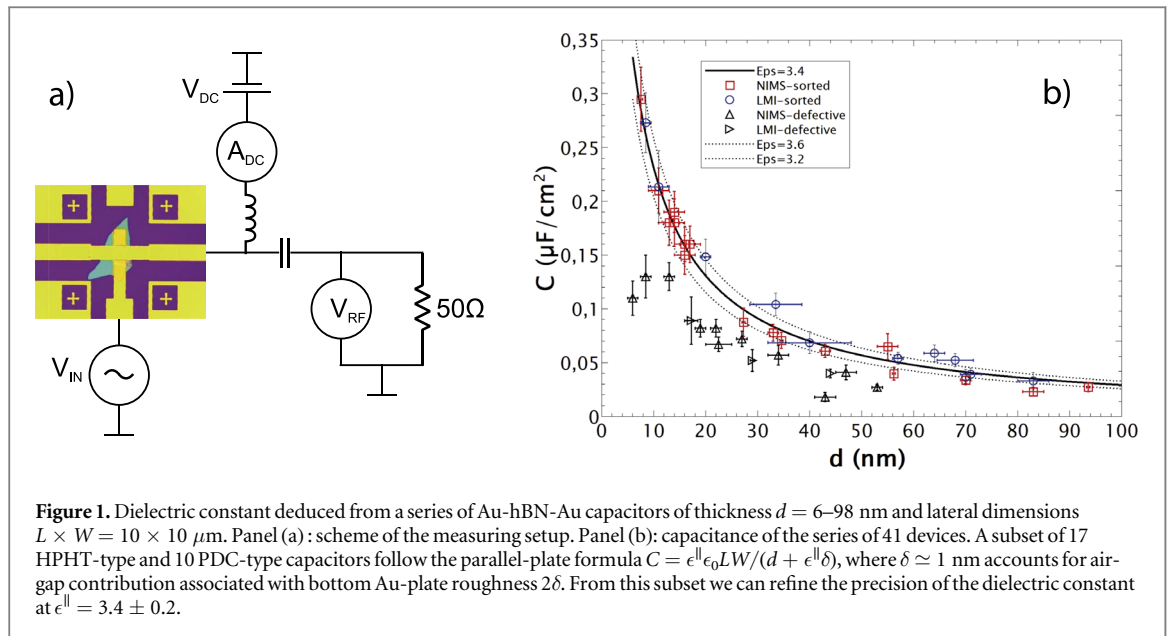
$\epsilon^{\parallel} \simeq 3.4 \pm 0.2$. The other 14 samples deviate from this law with lower capacitance values, presumably due to process imperfections involving spurious air gaps, due to dust or bubbles between the hBN flake and the bottom electrode. Besides, we do not see significant differences between HPHT and PDC crystals in terms of permittivity.

The dielectric strength is characterized by monitoring the leakage current at high bias which is analyzed in terms of a bulk conductivity. This analysis is carried out on a subset of 7 capacitors having survived the high-bias training. As shown in [9], dielectric conductivity is a three step process, starting by a threshold-less exponential current growth, followed by a quasi-saturation at a breakdown field E_{BD} and culminating by an irreversible current runaway for $E \gtrsim E_{BD}$ usually leading to sample breakdown. Here we focus on the pre-breakdown regime $E \lesssim E_{BD}$ where moderate current densities ($J < J_{BD} \simeq 0.5 \text{ A cm}^{-2}$) are applied that secure device integrity. In these conditions, we observe a bias-reversible, reproducible and polarity-independent behavior. The leakage current obeys a standard exponential growth with voltage which precludes unambiguous determination of a dielectric breakdown voltage (see figure 2(a)). However, when using an arbitrary breakdown current criterion $J \lesssim J_{BD}$, we find an overall increase of the breakdown voltage with thickness which suggests the relevance of a breakdown field, i.e. a bulk scenario, and justifies our breakdown analysis in terms of conductivity, e.g. $J/E(E)$ below a breakdown conductivity $\sigma_{BD} \sim 1 \mu\Omega \text{ cm}^{-1}$. We find that the conductivity, $\sigma = J/E$ obeys the Frenkel-Pool (FP) law (equation (1) below), corresponding to a trap-assisted, thermally-activated, bulk Schottky transport [14, 15]. Its signature lies in the linear dependence, $\ln \sigma / \sigma_{BD} = f(\sqrt{E/\epsilon^{\parallel} T^2})$ (see figure 2(b)), observed in the $[10^{-4} - 10^{-1}] \mu\text{S cm}^{-1}$ range at room temperature ($T = 300 \text{ K}$). The FP activation scenario is confirmed by the temperature dependence measured in one representative sample (inset of figure 2(b)). The robustness of the field dependence contrasts with the large variability (within a factor 10^6) of the conductivity prefactor. We assign the latter to a variability in the deep-level donor energy $\Phi_B \simeq 0.9 - 1.3 \text{ eV}$, with a logarithmical precision in the σ_{BD} prefactor. Remarkably, five devices show very similar Frenkel-Pool plots with deep-level traps energy $\Phi_{B,hBN} \simeq 1.27 \pm 0.03 \text{ eV}$, suggesting the existence of a quasi-intrinsic limit. This value is quite typical of that of technology-relevant insulators, such as Si_3N_4 where $\Phi_B \simeq 1.3 \pm 0.2 \text{ eV}$ [16], or SiO_2 where $\Phi_B \simeq 1 \text{ eV}$ [17] are reported. Some PDC-grown devices show however traps with a smaller energy, which opens routes for improvement of this less mature growth process.

2. Capacitor fabrication and setup

We have fabricated 41 Au-hBN-Au capacitors, 27 of them with HPHT-hBN from NIMS and 14 with the PDC-hBN from LMI. The growth technics, as well as structural and optical characterizations, of these high-quality crystals are detailed in [12] and [13, 18] respectively. The capacitors were deposited on high-resistivity Si substrates, suitable for RF measurement, that are covered with a 285 nm-thick SiO_2 layer. The bottom electrode and the RF coplanar waveguide structure are first deposited using laser lithography and thermal evaporation. The bottom 110/5 nm Au/Cr electrode is buried into SiO_2 to ensure a planar surface for hBN-flakes transfer. Planarisation is completed by mechanically polishing the small metallic pitches at the gold edges with isopropyl alcohol (IPA). This process minimizes air gaps between hBN and the bottom gold electrode which are ultimately limited by Au roughness of amplitude $2\delta \simeq 1.5 \text{ nm}$, as measured by atomic force microscopy (AFM). hBN crystals were mechanically exfoliated with polydimethylsiloxane (PDMS). We use poly-propylene carbonate PPC-PDMS stamps to dry transfer the hBN flakes on the bottom electrode. A second lithography step allows covering the hBN flake with a second Au/Cr electrode, which is conformal to the hBN dielectric and therefore airgap-free. The obtained structures form a nominally $10 \times 10 \mu\text{m}$ capacitor in a parallel-plate configuration. After annealing (1h at $240^\circ \text{ Celsius}$ under N_2) the capacitor's hBN thickness is determined by AFM and falls in the range $d = 6 - 98 \text{ nm}$. This range exceeds the minimal thickness (10–20 nm) for mobility-preserving encapsulation, is relevant for gate dielectric applications, and reaches the value ($\sim 100 \text{ nm}$) for fully-developed radiative cooling [19, 20]. It is however not relevant for tunnel-barrier applications which are described elsewhere [3–5]. The lateral size of capacitors was deliberately maximized at the limits of our exfoliation technique to increase experimental resolution, minimize spurious edge effects, and address the homogenous properties of hBN. It is precisely measured by scanning electron microscopy. We have indifferently used colinear and perpendicular source and drain electrodes, the later geometry being shown in the optical image of figure 1(a).

High frequency admittance measurements were carried out in a Janis (cryogenic) probe station under vacuum at room temperature (see sketch in figure 1(a)). The two-port scattering parameters S_{ij} of the capacitor were measured using an Anritsu MS4644B vectorial network analyzer (VNA) in the 10 MHz–10 GHz range. A short-open-load-reciprocal protocol was used to calibrate the wave propagation until the probe tips. As explained in [21], the wave propagation in the coplanar access of the capacitor is de-embedded by calculating the ABCD matrix from the S parameters of a symmetric thru-line reference structure. To correct for residual



parasitic stray capacitance effects, we convert the previous ABCD matrices into complex admittance matrices and subtract the contribution of a dummy reference structure of same geometry but devoid of the central Au-hBN-Au capacitor. The total capacitance is directly deduced from the low frequency (sub-GHz) imaginary part $Y_{12} = j\omega C$.

Breakdown measurements have been carried out on the same setup by monitoring the leakage current as function of the applied DC voltage (Keithley 2400 Source-Measure Unit). The same procedure has been applied at low temperature on a representative capacitor to check the activation mechanism at stake in the dielectric breakdown.

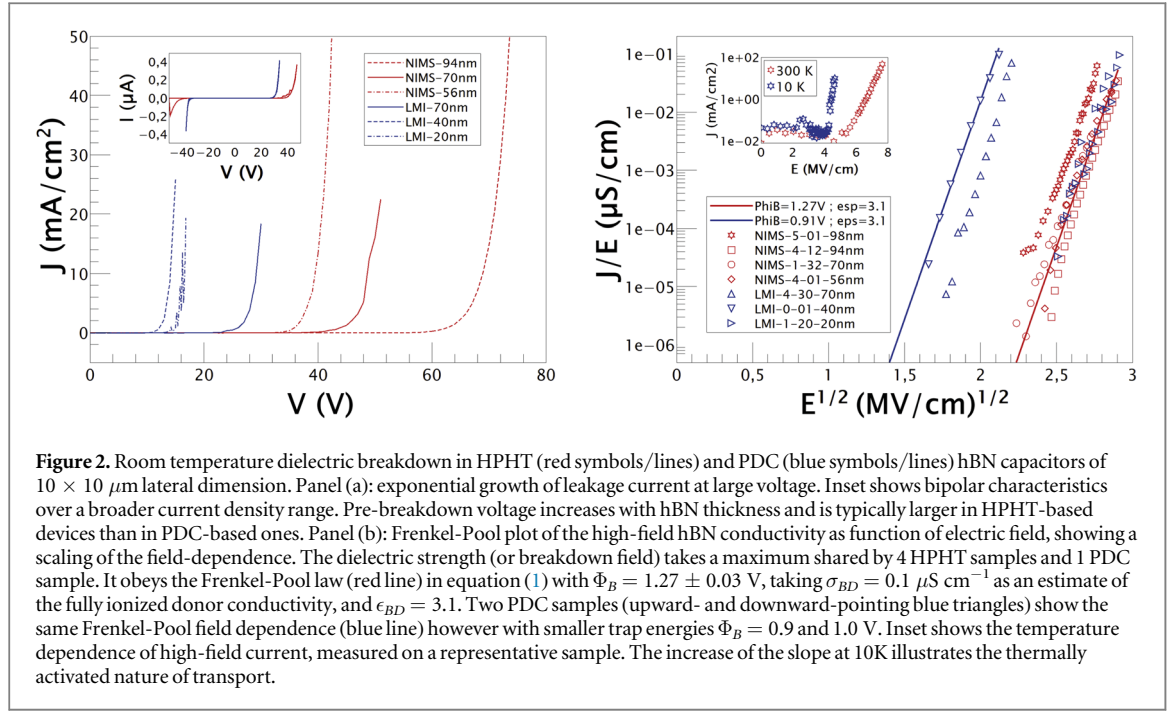
3. Dielectric constant

Figure 1(b) shows capacitances of the full set of 41 tested capacitors versus hBN thickness. Capacitance data, deduced from VNA measurements, are first compared with complementary data obtained by sub-MHz Lock-In techniques (not shown) to ascertain their frequency independence. Data are scattered, but we can still find a significant fraction of devices, of both HPHT and PDC sources, showing capacitance-data accumulation along an upper limit given by the parallel-plate capacitance formula $C = \epsilon_0 \epsilon^{\parallel} LW / d^*$ (red and blue symbols in figure 1(b)), where $\epsilon^{\parallel} \simeq 3.4 \pm 0.2$ and $d^* = d + \epsilon^{\parallel} \delta$ is an effective dielectric thickness accounting for spurious air-gap contributions due to (bottom) metal roughness $2\delta \simeq 2$ nm. One third of the series (14 devices, black symbols in figure 1(b)), exhibiting lower capacitance values caused by process imperfections including spurious air gaps, presumably due to dust or bubbles between the hBN flake and the bottom electrode, are discarded. From the selected devices (17 HPHTs and 11 PDCs), we are able to narrow down the dielectric constant window and provide a recommended value of the hBN dielectric constant $\epsilon^{\parallel} = 3.4 \pm 0.2$. This value exceeds by 13% the $\epsilon^{\parallel} \simeq 3.0$ reported in metal-hBN-graphene quantum-Hall devices [8, 19]. This apparent disagreement can be lifted when considering the series quantum capacitance of the graphene electrode in these thin hBN samples [8, 22], as well as roughness-induced air-gaps at the bottom electrodes. Our measured permittivity turns out to be consistent with the theoretical prediction $\epsilon_{\parallel} = 3.38$ of [23], following an ab-initio approach which already quantitatively predicts correctly the optical permittivity and its anisotropy [24].

4. Breakdown current and conductivity

Figure 2(a) shows typical current-voltage characteristics of capacitors showing a strongly non-linear onset of current. Characteristics become irreversible at large current density, $J \sim 0.1$ $\mu\text{A cm}^{-2}$, eventually leading to sample breakdown for $J \gtrsim J_{BD} \gtrsim 1$ $\mu\text{A cm}^{-2}$. At the lower bias of figure 2(a), I-V curves are reproducible and bias symmetric (inset of figure 2(a)). As seen in the figure, the breakdown voltage V_{BD} shows a tendency to increase with hBN thickness d , suggesting a bulk origin of breakdown.

A deeper insight into the breakdown mechanism is provided by figure 2(b) which reveals a $\ln(J/E) \propto \sqrt{E}$ scaling of conductivity. It corresponds to the Frenkel-Pool (FP) effect [14–16], where a finite dielectric



conductivity J/E arises due to thermal de-trapping of deep-level electrons of energy $e\Phi_B \gg k_B T$. Conductivity increases at large voltage as electric field lowers the barrier height Φ_B by an amount $\sqrt{eE/\pi\epsilon_{BD}\epsilon_0}$. At ultimate fields, it eventually leads to full ionization of traps whenever $(\Phi_B - \sqrt{eE/\pi\epsilon_{BD}\epsilon_0}) \lesssim k_B T$, which defines a breakdown electric field E_{BD} . In this picture, $E_{BD} = \Phi_B/r_0$ is on the order of the impurity field, where $r_0 = e/\pi\epsilon_{BD}\epsilon_0\Phi_B$ is the screening length. The FP conductivity writes [15]:

$$\frac{J}{E} = \sigma_{BD} \times \exp\left[-e\frac{\Phi_B - \sqrt{eE/\pi\epsilon_{BD}\epsilon_0}}{k_B T}\right], \quad (1)$$

where the breakdown conductivity $\sigma_{BD} = J_{BD}/E_{BD} \simeq N_T e \mu$ corresponds to the band conductivity (mobility μ) for fully ionized traps (concentration N_T). Equation (1) holds in the field range $E_{FN} \lesssim E \lesssim E_{BD}$, or $J = 10^{-4} - 10^{-1} \mu\text{A cm}^{-2}$, where E_{FN} is the Fowler-Nordheim tunneling limit for a defect-free thin triangular barrier [16].

As seen in figure 2(b), the field dependence of breakdown conductivity does obey the FP mechanism for both hBN sources, with a slope solely determined by ϵ_{BD} . Data fitting suggests a small field-suppression of permittivity with $\epsilon_{BD} \simeq 3.1 < \epsilon^{\parallel} \simeq 3.4$. In contrast to the universality of slope, the prefactor $\sigma_{BD} \exp\left(-\frac{e\Phi_B}{k_B T}\right)$ shows sample variability, exhibiting variations by six orders of magnitude. This observation highlights the strong sensitivity of breakdown to material quality which, in the FP scenario, mainly stems from the variability in the trap potential Φ_B (and E_{BD}), as that of N_T and μ in σ_{BD} cannot explain such a large scatter alone. For a quantitative estimation of Φ_B we set $\sigma_{BD} \sim 0.1 \mu\text{S cm}^{-1}$, corresponding to $J_{BD} \sim 0.5 \text{ A cm}^{-2}$ and $E_{BD} \sim 5 \text{ MV cm}^{-1}$. This breakdown current is typical of quasi saturation values observed in our samples, and in the literature (figure 5 in [9]). With this procedure, we extract $\Phi_B = 0.9 - 1.3 \text{ V}$ in figure 2(b). Remarkably, we observe an accumulation of conductivity data (4 HPHT and 1 PDC capacitors) along an upper limit represented by the $\Phi_B = 1.27 \pm 0.03 \text{ V}$ line. This suggests the existence of a dielectric strength limit in high-quality hBN crystals. Dispersion among that data-subset of highest-quality samples corresponds to $\Delta\Phi_B = 0.06 \text{ V}$, or a variation of σ_{BD} in the range $0.01 - 1 \mu\text{S cm}^{-1}$. Taking a typical insulator mobility $\mu \lesssim 1 \text{ cm}^2/\text{Vs}$, this translates into a trap density $N_T \sim 10^{12} - 10^{13} \text{ cm}^{-3}$ and a trap number $N_T dLW \sim 10 - 100$.

To further establish the FP mechanism of breakdown we have added, in the insert of figure 2(b), a comparison between 10 K and 300 K breakdown current measurements performed on an additional sample (NIMS-5-01-98 nm), which illustrates the strong temperature dependence of activated FP transport, at variance with tunneling-based scenarios.

5. Conclusion

Our estimate of hBN dielectric permittivity, $\epsilon^{\parallel} \equiv \epsilon_0^{\parallel c} = 3.4 \pm 0.2$, agrees with the calculations of Ohba *et al*, which predicts: $\epsilon_{\infty}^{\perp c} = 4.85$, $\epsilon_{\infty}^{\parallel c} = 2.84$, $\epsilon_0^{\perp c} = 6.61$, $\epsilon_0^{\parallel c} = 3.38$ [23]. This completes previous results based on optical measurements giving $\epsilon_{\infty}^{\perp c} = 4.95$, $\epsilon_{\infty}^{\parallel c} = 2.86$, $\epsilon_0^{\perp c} = 6.96$ [24]. The excellent agreement in the four relevant dielectric constants of hBN gives strong confidence in the ab-initio technique to provide reliable predictions of static and dynamical properties of BN crystals including those of the zinc-blende and wurtzite crystals.

The relevance of the 3D Frenkel-Pool mechanism of conductivity in the 2D hBN was not granted as 2D materials may sustain specific mechanisms. We demonstrate here that it works for *c*-axis transport, but the situation can be different for in-plane electric fields with the opening of new leakage channels associated with charges gliding in-between hBN planes. The identification of deep-level traps responsible for breakdown is beyond the scope of our work, especially as leakage current alone cannot identify the acceptor/donor nature of the levels. We find $\Phi_B \simeq 1.27 \pm 0.03$ eV in hBN, which is larger than the 1 eV reported in SiO₂ [17], and comparable with the 1.3 ± 0.2 eV in Si₃N₄ [16]. In the latter case, deep traps are attributed to silicon-dangling-bond centers [25]. This trap energy determines a maximum breakdown field $E_{BD} \simeq 5$ MV cm⁻¹, defined as the threshold for current quasi-saturation.

Finally we conclude on the strong similarity of HTHP and PDC hBN crystals in terms of dielectric permittivity and strength, with however a better yield in terms of intrinsic dielectric breakdown for the NIMS samples, which can to a large extent be attributed to a longer maturity of the growth technique.

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Data availability statement

The data that support the findings of this study are available upon reasonable request from the authors.

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